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AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions and listings of claims in the application:

Claim 1 (Currently amended): A method for the prophylaxis or treatment of a HSV-1, HSV-2, or CMV infection in a subject, comprising administering to a subject in need of such treatment a therapeutically effective amount of at teast one pharmacologically acceptable oligonucleotide at teast 3029 nucleotides in length, wherein the anti-viral activity of said oligonucleotide occurs principally by a non-sequence complementary mode of action, and wherein said oligonucleotide comprises at least one phosphorothioated linkage.

Claim 2 (Original): The method of claim 1, wherein said subject is a human.

Claim 3-13 (Canceled).

Claim 14 (Currently amended): The method, pharmaceutical composition, or kit of claim 1, or 23, or 12, wherein said at least one antiviral oligonucleotide comprises at least one antiviral randomer oligonucleotide.

Claim 15 (Currently amended): The method, pharmaceutical composition, or kit of claim 1, or 23, or 12, wherein said oligonucleotide is not complementary to any portion of the genomic sequence of HSV-1, HSV-2, or CMV.

Claim 16 (Canceled).

Claim 17 (Currently amended): The method, pharmaceutical composition, or kit of claim 1, or 23, or 12, wherein said oligonucleotide is at least 40 nucleotides in length.

Claim 18 (Currently amended): The method, pharmaceutical composition, or kit of claim 1 or, 23, or 12, wherein each said oligonucleotide comprises at least one modification to its chemical structure.

Claim 19-20 (Cancelled).

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Claim 21 (Currently amended): The method, pharmaceutical composition, or kit of claim 1 or, 23, or 12, wherein each said oligonucleotide comprises at least one 2'-modification to the ribose moiety.

Claim 22 (Currently amended): The method, pharmaceutical composition, or kit of claim 1 or, 23, or 12, wherein each said oligonucleotide comprises at least one methylphosphonate linkage.

Claim 23-26 (Cancelled).

Claim 27 (Currently amended): The method, pharmaceutical composition, or kit of claim 1 or, 23, or 12, wherein said oligonucleotide is double stranded.

Claim 28 (Currently amended): The method, pharmaceutical composition, or-kit of claim 1 or, 23, or 12, wherein said oligonucleotide binds to one or more viral components.

Claim 29 (Currently amended): The method, pharmaceutical composition, or kit of claim 1_or, 23, or 12, wherein at least a portion of the sequence of said oligonucleotide is derived from a viral genome.

Claim 30-38 (Cancelled).

Claim 39 (New): The method of claim 1 or 2, wherein each nucleotide of said oligonucleotide are linked to one another by a phosphorothioated linkage.

Claim 40 (New): The method of claim 1 or 2, wherein said oligonucleotide is selected from the group consisting of REP 2005, REP 2006, REP 2007, REP 2008, SEQ ID NO: 6, SEQ ID NO: 9, REP 2024, SEQ ID NO: 20, SEQ ID NO: 23, SEQ ID NO: 25, SEQ ID NO: 26 and REP 2060.

Claim 41 (New): The method of claim 1 or 2, wherein said oligonucleotide is SEQ ID NO: 22.

Claim 42 (New): The method of claim 1 or 2, wherein said oligonucleotide is SEQ ID NO: 24.